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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/750,846	01/05/2004	Heimo Haikala	06267.0116	3865
22852 7	590 01/27/2006		EXAMINER	
FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER LLP 901 NEW YORK AVENUE, NW WASHINGTON, DC 20001-4413			HENLEY III, RAYMOND J	
			ART UNIT	PAPER NUMBER
			1614	

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Please find below and/or attached an Office communication concerning this application or proceeding.

•	Application No.	Applicant(s)			
	10/750,846	HAIKALA ET AL.			
Office Action Summary	Examiner	Art Unit			
	Raymond J. Henley III	1614			
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply					
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.  - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).					
Status					
1)☐ Responsive to communication(s) filed on  2a)☐ This action is FINAL. 2b)☒ This  3)☐ Since this application is in condition for allowar closed in accordance with the practice under E	action is non-final. nce except for formal matters, pro				
Disposition of Claims					
4) ☐ Claim(s) 3 and 4 is/are pending in the application 4a) Of the above claim(s) is/are withdraw 5) ☐ Claim(s) is/are allowed.  6) ☐ Claim(s) 3 and 4 is/are rejected.  7) ☐ Claim(s) is/are objected to.  8) ☐ Claim(s) are subject to restriction and/or	wn from consideration.	·			
Application Papers					
9)☐ The specification is objected to by the Examine 10)☒ The drawing(s) filed on <u>05 January 2004</u> is/are: Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11)☐ The oath or declaration is objected to by the Ex	: a) $\square$ accepted or b) $\square$ objected drawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119					
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).  a) ■ All b) ■ Some * c) ■ None of:  1. ■ Certified copies of the priority documents have been received.  2. ■ Certified copies of the priority documents have been received in Application No. ■  3. ■ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).  * See the attached detailed Office action for a list of the certified copies not received.					
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)  Paper No(s)/Mail Date 1/5/04.	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:				

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## **CLAIMS 3 AND 4 ARE PRESENTED FOR EXAMINATION**

Applicants' Preliminary Amendment and Information Disclosure Statement filed January 5, 2004 have been received and entered into the application. Accordingly, claims 1-2 have been canceled and the specification at page 1 has been amended. Also, as reflected by the attached, completed copy of form PTO-1449, the Examiner has considered the cited references.

#### Claim Objection

Claim 3 is objected to as being incomplete. In particular, claim 3 merely reads that mortality may be reduced in a mammal with congestive heart failure, but fails to specify that such mortality would be due to the heart failure. In order to overcome this objection, it is suggested that claim 3 be amended to indicate that the mortality is due to congestive heart failure.

### Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later

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invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

The conclusion of obviousness based on the remaining two references is proper because Haikala et al. (U.S. Patent No. 5,905,078) and Sircar (U.S. Patent No. 4,397,854) would be proper references under 35 U.S.C. § 102(b).

Claims 3-4 are rejected under 35 U.S.C. 103(a) as being obvious over Haikala et al. (U.S. RE38,102, "Haikala '102"), Haikala et al. (U.S. Patent No. 5,905,078; "Haikala et al. '078") or Applicants' acknowledgment at page 1, lines 2-4 of the third paragraph, or Sircar (U.S. Patent No. 4,397,854) in view of Campbell (U.S. Patent No. 4,432,979) and Diamond et al. (U.S. Patent No. 4,517,310).

The applied references, i.e., Haikala et al. (U.S. RE38,102) has a common assignee and inventor with the instant application. Based upon the earlier effective U.S. filing date of the reference, it constitutes prior art only under 35 U.S.C. 102(e). This rejection under 35 U.S.C. 103(a) might be overcome by: (1) a showing under 37 CFR 1.132 that any invention disclosed but not claimed in the reference was derived from the inventor of this application and is thus not an invention "by another"; (2) a showing of a date of invention for the claimed subject matter of the application which corresponds to subject matter disclosed but not claimed in the reference, prior to the effective U.S. filing date of the reference under 37 CFR 1.131; or (3) an oath or declaration under 37 CFR 1.130 stating that the application and reference are currently owned by the same party and that the inventor named in the application is the prior inventor under 35 U.S.C. 104, together with a terminal disclaimer in accordance with 37 CFR 1.321(c). This rejection might also be overcome by showing that the reference is disqualified under 35 U.S.C.

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103(c) as prior art in a rejection under 35 U.S.C. 103(a). See MPEP § 706.02(l)(1) and § 706.02(l)(2).

Haikala '102, Haikala et al. '078" or Sircar (U.S. Patent No. 4,397,854) teach racemic (Sicar) or R-N-[4-(1,4,5,6-tetrahydro-4-methyl-6-oxo-3-pyridazinyl)phenyl]acetamide as being effective cardiotonic agents which are effective for the treatment or prevention of congestive heart failure. See Haikala '102 at the abstract, col. 1, line 13, 14, 16, and 17; Haikala et al. '078 at the abstract, col. 1, lines 8, 9, 11 and 12; and Sircar (teaches cardiotonic activity only at col. 1, lines 12-14; col. 6, lines 21-26 and claim 1 which encompasses the presently claimed acetamide compound. Also, Applicants' acknowledge at page 1 of the specification, lines 3-4 of the second paragraph that the acetamide compound was known for the treatment of chronic heart failure.

The difference between the above and the claimed subject matter lies in that none of the above references teach that mortality reduction, (presumably from heart failure), may be achieved through administration of the claimed acetamide compound.

However, the difference between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains because the artisan would have appreciated that if a compound is effective against a disease which can cause mortality, the incidence of mortality should decrease. The basis of this position rests upon a logical conclusion.

It is noted that Sicar does not highlight congestive heart failure, *per se*, however, it was well known in the art that cardiotonic agents were effective for the treatment of heart failure, (see Campbell and Diamond et al. at the abstract).

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#### **Double Patenting**

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., In re Berg, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); In re Goodman, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); In re Longi, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); In re Van Ornum, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); In re Vogel, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and In re Thorington, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claims 3-4 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 10 of U.S. Patent No. Poder et al. (U.S. Patent No. 6,949,548). Although the conflicting claims are not identical, they are not patentably distinct from each other because the difference between the patented claim and the present claim lies in that the patented claim fails to highlight the administration of the R-isomer of the acetamide compound as claimed. That is, the patented claims teaches levosimendan for the presently claimed purpose of reducing mortality. The patented claim contains all of the remaining elements of the present claims.

However, the difference between the patented claim and present claims would have been obvious to one of ordinary skill in the art because, as acknowledged by Applicants at page 1, final, partial paragraph of the present specification, the claimed R isomer is present as an active

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metabolite in humans following administration of levosimendan. Accordingly, the patented claim inherently teaches that which is presently claimed.

Regarding the legal standard for anticipation/inherency Under - 35 USC § 102, it is well settled that to anticipate a claim under 35 U.S.C. § 102, a single prior art reference must place the invention in the public's possession by disclosing each and every element of the claimed invention in a manner sufficient to enable one skilled in the art to practice the invention. Scripps Clinic & Research Foundation v. Genetech, Inc., 927 F.2d 1565, 1576, 18 U.S.P.Q.2d 1001, 1001 (Fed. Cir. 1991); In re Donahue, 766 F2d531, 533, 226 U.S.P.Q. 619, 621 (Fed. Cir. 1985). To anticipate, the prior art must either expressly or inherently disclose every limitation of the claimed invention. MEHL/Biophile Int'l Corp. v. Milgraum, 192 F.3d 1362, 1365, 52 U.S.P.Q.2d 1303, 1303 (Fed. Cir. 1999) (citing to In re Schreiber, 128 F.3d 1473, 1477, 44 U.S.P.Q. 1429, 1431 (Fed. Cir. 1997)); Atlas Powder Co. v. Ireco Inc., 190 F.3d 1342, 1347, 51 U.S.P.Q.2d 1943, 1946 (Fed. Cir. 1999). Additionally, and most importantly, to inherently anticipate, the prior art must necessarily function in accordance with, or include, the claimed limitations. MEHL/Biophile, 192 F.3d at 1365, 52 U.S.P.Q.2d at 1303. However, it is not required that those of ordinary skill in the art recognize the inherent characteristics or the function of the prior art. Id. Specifically, discovery of the mechanism underlying a known process does not make it patentable.

Accordingly, for the above reasons, the claims are deemed properly rejected and none are allowed.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Raymond J. Henley III whose telephone number is 571-272-0575. The examiner can normally be reached on M-F, 8:30 am to 4:00 pm Eastern Time.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christopher Low can be reached on 571-272-0951. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Raymond J Henley III Primary Examiner Art Unit 1614

January 24, 2006